## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

- (Original) A method for inhibiting the growth and/or spreading of malignant tumors, metastases and/or lung foci, comprising administering a composition comprising Nα(2,4,6-Triisopropylphenylsulfonyl)-3-amidino-(D,L)-phenylalanine 4-ethoxycarbonylpiperazide, the L enantiomer thereof or a pharmaceutically suitable salt thereof and a pharmaceutically acceptable carrier to a patient in need of such inhibition.
- 2. (Original) The method according to claim 1, wherein said tumor affects lymphatic tissue.
- 3. (Original) The method according to claim 2, wherein said lymphatic tissue is lymph nodes.
- 4. (Original) The method according to claim 3, wherein said lymph nodes are selected from the group consisting of axillary lymph nodes and intraperitoneal lymph nodes.
- 5. (Original) The method according to claim 1, further comprising administering a cytotoxic substance.

- 6. (Previously presented) The method according to claim 5, wherein said cytotoxic substance is selected from the group consisting of cisplatin, carboplatin, doxorubicin, epirubicin, 5-fluorouracil and a taxane.
- 7. (Original) The method according to claim 6, wherein said taxane is paclitaxel.
- 8. (Original) The method according to claim 1, wherein said malignant tumors are breast cancer.
- 9. (Original) The method according to claim 1, wherein said composition is administered once daily to once weekly.
- 10. (Original) A pharmaceutical composition comprising Nα(2,4,6-Triisopropylphenylsulfonyl)-3-amidino-(D,L)-phenylalanine 4-ethoxycarbonylpiperazide, the L enantiomer thereof or a pharmaceutically suitable salt thereof, an additional, pharmacologically active substance, and a pharmaceutically acceptable carrier.
- 11. (Original) The pharmaceutical composition according to claim 10, wherein said additional, pharmacologically active substance is selected from the group consisting of radio labels or cytotoxic substances.
- 12. (Original) The pharmaceutical composition according to claim 11, wherein said additional, pharmacologically active substance is a cytotoxic substance.

- 13. (Previously presented) The pharmaceutical composition according to claim 12, wherein said cytotoxic substance is selected from the group consisting of cisplatin, carboplatin, doxorubicin, epirubicin, 5-fluorouracil and a taxane.
- 14. (Original) The pharmaceutical composition according to claim 13, wherein said taxane is paclitaxel.
- 15. (Original) A kit comprising, in separate containers, a) Nα(2,4,6-Triisopropylphenylsulfonyl)-3-amidino-(D,L)-phenylalanine 4-ethoxycarbonylpiperazide, the L enantiomer thereof or a pharmaceutically suitable salt thereof, and b) radio labels and/or cytotoxic substances.
- 16. (Original) A method for the treatment of malignant tumors comprising
- a) surgically removing a primary tumor from a patient, and
- b) administering a composition comprising Nα(2,4,6-Triisopropylphenylsulfonyl)-3-amidino-(D,L)-phenylalanine 4-ethoxycarbonylpiperazide, the L enantiomer thereof or a pharmaceutically suitable salt thereof and a pharmaceutically acceptable carrier to said patient.
- 17. (Original) The method according to claim 16, further comprising administering cytotoxic agents and/or radiation therapy to said patient.

- 18. (New) A method for inhibiting the growth and/or spreading of malignant tumors, metastases and/or lung foci, comprising contacting a cell with  $N\alpha(2,4,6-Triisopropylphenylsulfonyl)$ -3-amidino-(D,L)-phenylalanine 4-ethoxycarbonylpiperazide, the L enantiomer thereof or a pharmaceutically suitable salt thereof, wherein said cell is in a patient in need of such inhibition.
- 19. (New) A method for inhibiting the growth and/or spreading of malignant tumor cells metastases and/or lung foci in a patient in need of such inhibition, comprising administering  $N\alpha(2,4,6\text{-Triisopropylphenylsulfonyl})$ -3-amidino-(D,L)-phenylalanine 4-ethoxycarbonylpiperazide, the L enantiomer thereof or a pharmaceutically suitable salt thereof to a tumor cell in said patient.